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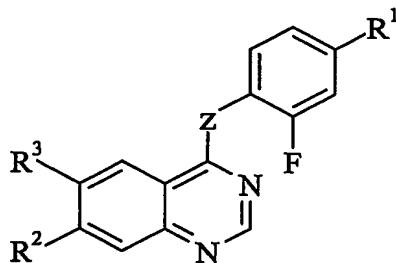
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(54) Title: QUINAZOLINE DERIVATIVES AS INHIBITORS OF VEGF RECEPTOR TYROSINE KINASES



(I)

(57) Abstract: The present invention relates to compounds of the Formula (I): wherein Z is -NH-, -O- or -S-; R<sup>1</sup> represents bromo or chloro; R<sup>3</sup> represents C<sub>1-3</sub> alkoxy or hydrogen; R<sup>2</sup> is selected from one of the following three groups: (i) Q<sup>1</sup>X<sup>1</sup> - wherein X<sup>1</sup> and Q<sup>1</sup> are as defined herein; (ii) Q<sup>15</sup>W<sup>3</sup> - wherein Q<sup>15</sup> and W<sup>3</sup> are as defined herein; and (iii) Q<sup>21</sup>W<sup>4</sup>C<sub>1-3</sub>alkylX<sup>1</sup> wherein X<sup>1</sup>, W<sup>4</sup> and Q<sup>21</sup> are as defined herein; and salts thereof; their use in the manufacture of a medicament for use in the production of an antiangiogenic and/or vascular permeability reducing effect in warm blooded animals; processes for the preparation of such compounds; pharmaceutical compositions containing a compound of formula (I) or a pharmaceutically acceptable salt thereof and methods of treating disease states involving angiogenesis by administering a compound of formula (I) or a pharmaceutically acceptable salt thereof. The compounds of formula (I) inhibit the effects of VEGF, a property of value in the treatment of a number of disease states including cancer and rheumatoid arthritis.

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SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ,  
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